Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

- 1. (Currently amended) A An isolated and purified polypeptide comprising an enzyme activity, wherein the enzyme activity the following physicochemical properties (1) to (5):(1)It asymmetrically reduces N-benzyl-3-pyrrolidinone to produce (S)-N-benzyl-3-pyrrolidinol with NADPH as a coenzyme;(2)Optimum, wherein the enzyme activity has an optimum action pH: of 4.5 to 5.5;(3)—, an optimum Optimum action temperature: of 40°C to 45°C;(4) and, a molecular Molecular weight: of Aabout 29,000 as determined by gel filtration analysis and about 35,000 as determined by SDS-polyacrylamide gel electrophoresis analysis; and(5) Inhibitor: wherein the enzyme activity It-is inhibited by the divalent copper ion.
- 2. (Currently amended) A An isolated and purified polypeptide comprising described in the following (a) or (b):
- (a) A polypeptide having the amino acid sequence shown under SEQ ID NO: 1 in the sequence listing; or
- (b) A polypeptide having an amino acid sequence obtainable derived from the amino acid sequence shown under SEQ ID NO: 1 in the sequence listing by substitution, insertion, deletion and/or addition of one or more amino acids, wherein the polypeptide possesses and having an enzyme activity in comprising asymmetrically reducing N-benzyl-3-pyrrolidinone to produce (S)-N-benzyl-3-pyrrolidinol.
- 3. (Currently amended) The polypeptide according to Claim of claim 1, wherein the polypeptide which is derived from a microorganism belonging to the genus Micrococcus.
- 4. (Currently amended) The polypeptide according to Claim of claim 3, wherein said the microorganism is the strain Micrococcus luteus Micrococcus luteus IFO 13867.
- 5. (Currently amended) An isolated and purified DNA molecule coding for the polypeptide according to Claim of claim 1.

- 6. (Currently amended) An isolated and purified DNA molecule coding for a polypeptide having wherein the polypeptide comprises an enzyme activity in comprising asymmetrically reducing N-benzyl-3-pyrrolidinone to produce (S)-N-benzyl-3-pyrrolidinol, and wherein the DNA molecule hybridizing hybridizes with a DNA having to a-nucleotide comprising sequence shown under of SEQ ID NO: 2 in the sequence listing under stringent conditions.
- 7. (Currently amended) An isolated and purified DNA molecule coding for a polypeptide, wherein the polypeptide possesses having enzyme activity in comprising asymmetrically reducing N-benzyl-3-pyrrolidinone to produce (S)-N-benzyl-3-pyrrolidinol, and wherein the sequence of the DNA molecule has having at least 60% sequence identity with a nucleotide sequence shown under to SEQ ID NO: 2 in the sequence listing.
- 8. (Currently Amended) An expression vector eontaining comprising the isolated DNAs molecule according to Claim 1 of claim 5.
- 9. (Currently amended) The expression vector according to Claim of claim 8, wherein the vector which is a plasmid pTSBH.
- 10. (Currently amended) The expression vector according to Claim of claim 8, wherein the isolated DNA molecule codes which contains a DNA coding for a polypeptide having glucose dehydrogenase activity.
- 11. (Currently amended) The expression vector according to Claim of claim 10, wherein said the polypeptide having glucose dehydrogenase activity is a <u>Bacillus megaterium</u>. Bacillus megaterium derived glucose dehydrogenase.
- 12. (Currently amended) The expression vector according to Claim of claim 11, wherein the vector which is a plasmid pTSBG1.
- 13. (Currently Amended) A transformant containing comprising the expression vector according to Claim 1 of claim 8.
- 14. (Currently Amended) A transformant containing both the expression vector according to Claim 1 of claim 8 and an expression vector containing a DNA molecule coding for a polypeptide having glucose dehydrogenase activity.

- 15. (Currently amended) The transformant according to Claim of claim 14, wherein said the polypeptide having glucose dehydrogenase activity is a <u>Bacillus megaterium</u> <u>Bacillus megaterium</u> derived glucose dehydrogenase.
- 16. (Currently amended) The transformant according to Claim of claim 1 13, wherein a host thereof is Escherichia coli Escherichia coli.
- 17. (Currently amended) The transformant according to Claim of claim 16, wherein the host which is Escherichia coli Escherichia coli HB101 (pTSBH).
- 18. (Currently amended) The transformant according to Claim of claim 16, wherein the host which is Escherichia coli Escherichia coli HB101 (pTSBG1).
- 19. (Currently amended) The transformant according to Claim of claim 16, wherein the host which is Escherichia coli Escherichia coli HB101 (pTSBH, pSTVG).
- 20. (Currently Amended) A production method of producing (S)-N-benzyl-3-pyrrolidinol comprising:

a step of a) reacting the transformant according to Claim of claim 1 13 and/or a treated product thereof with N-benzyl-3-pyrrolidinone, and

a step of b) harvesting the thus produced (S)-N-benzyl-3-pyrrolidinol produced in a).

- 21. (Currently amended) The method according to Claim of claim 20, wherein the step of reacting is carried out in the presence of a coenzyme regenerating system.
 - 22. (New) An expression vector comprising the isolated DNA-molecule of claim 6.
 - 23. (New) An expression vector comprising the isolated DNA-molecule of claim 7.